

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.**

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : C07D 251/54, C09K 21/12, C08K 5/3492, C08G 79/04	A1	(11) International Publication Number: WO 00/02869 (43) International Publication Date: 20 January 2000 (20.01.00)
(21) International Application Number: PCT/NL99/00426 (22) International Filing Date: 7 July 1999 (07.07.99) (30) Priority Data: 1009588 8 July 1998 (08.07.98) NL (71) Applicant (for all designated States except US): DSM N.V. [NL/NL]; Het Overloon 1, NL-6411 TE Heerlen (NL). (72) Inventors; and (75) Inventors/Applicants (for US only): KERSJES, Johanna, Gertruda [NL/NL]; Gaarstraat 25, NL-6121 HG Born (NL). KIERKELS, Renier, Henricus, Maria [NL/NL]; Veldstraat 12, NL-6099 AT Heel (NL). (74) Agent: VAN BOKHOVEN, Petrus, Hubertus, Maria; Octrooibureau DSM, P.O. Box 9, NL-6160 MA Geleen (NL).		(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: POLYPHOSPHATE SALT OF A 1,3,5-TRIAZINE COMPOUND WITH A HIGH DEGREE OF CONDENSATION AND ITS USE AS FLAME RETARDANT IN POLYMER COMPOSITIONS		
(57) Abstract Salt of a 1,3,5-triazine compound with polyphosphoric acid with a virtually linear structure, with the number average degree of condensation n being higher than 20, the content of the 1,3,5-triazine compound being higher than 1.1 mole per mole of phosphorus atom and the pH of a 10 % slurry in water being higher than 4.5. Further, a process for the preparation of the polyphosphate salt of a 1,3,5-triazine compound with a number average degree of condensation n that is higher than 20, by converting a 1,3,5-triazine compound with orthophosphoric acid at room temperature into the phosphate of the 1,3,5-triazine compound, after which this salt is converted at elevated temperature into a polyphosphate of the 1,3,5-triazine compound, and further flame retardant polymer compositions comprising the polyphosphate salt of a 1,3,5-triazine compound with a number average degree of condensation n higher than 20.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

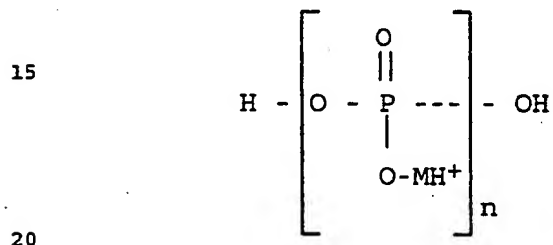
AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LJ	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

POLYPHOSPHATE SALT OF A 1,3,5-TRIAZINE COMPOUND WITH A HIGH
DEGREE OF CONDENSATION AND ITS USE AS FLAME RETARDANT IN
POLYMER COMPOSITIONS

5

The invention relates to a polyphosphate salt of a 1,3,5-triazine compound, a process for its preparation, and use of the resulting salt as flame
10 retardant in polymer compositions.

The polyphosphate of a 1,3,5-triazine compound can be represented by the general formula:



where M represents a 1,3,5-triazine compound and n is an integer greater than 3 that represents a measure of the number average degree of condensation. For high values of n, the polyphosphate of a 1,3,5-triazine
25 compound can best be represented by the formula (MHPO₃)_n. Theoretically, the structure is substantially linear if the M/P (triazine/phosphorus) ratio is almost exactly 1.0. Similarly, if the M/P ratio is less than 1, it indicates that the product includes some
30 crosslinking and, if the M/P ratio is less than 0.4, it indicates that the degree of crosslinking is sufficient for the product to form a network structure.

Melamine polyphosphate and a process for preparing melamine polyphosphate are described, inter
35 alia, in WO 97/44377. According to this reference,

melamine polyphosphate having a solubility of 0.01 to 0.10 g per 100 ml water at 25°C, a pH between 2.5 and 4.5, and a melamine/phosphorus molar ratio of between 1.0 and 1.1, may be obtained as a 10 wt% aqueous slurry at 25°C. WO 97/44377 also describes a two-step process for preparing the disclosed melamine polyphosphate slurry. In the first step melamine, urea, and an aqueous orthophosphoric acid solution (containing at least 40 wt% orthophosphoric acid), are mixed to produce a reaction mixture having a melamine/orthophosphoric acid molar ratio between 1.0 and 1.5 moles and a urea/orthophosphoric acid molar ratio between 0.1 and 1.5 at a temperature between 0 and 140°C. The resulting reaction mixture is then stirred at a temperature between 0 and 140°C and dehydrated to produce a powdery product comprising a double salt of orthophosphoric acid with melamine and urea. This powdery product is then heated to between 240 and 340°C and maintained in this temperature range for between 0.1 and 30 hours while preventing agglomeration to obtain melamine polyphosphate.

One disadvantage of melamine polyphosphates having a melamine/phosphorus molar ratio between 1.0 and 1.1 such as those prepared according to WO 97/44377 is their general unsuitability for use as a flame retardant in polymers. This is particularly the case for polymers such as nylons and polyesters that are typically processed at elevated temperatures, temperatures at which the salts do not exhibit sufficient thermal stability. Moreover, the pH of such salts are relatively low, a property that tends to adversely affect the polymer's mechanical properties such as impact strength, tensile strength, and breaking strength.

It has been found, however, that salts of 1,3,5-triazine compounds with polyphosphoric acid having n values greater than 20, and preferably greater than 40, and M/P ratios of at least 1.1, and preferably at least 1.2, do not exhibit these disadvantages when combined with polymers. Further, according to the present invention, the n value of such salts should generally be between 20 and 200, preferably between 40 and 150, and the M/P ratio should be between 1.1 and 2.0, preferably between 1.2 and 1.8. Further, the pH of a 10 wt% aqueous slurry of salts prepared according to the present invention will generally be greater than 4.5 and preferably at least 5.0. The referenced pH value is determined by introducing 25 g of the salt and 225 g of pure, 25°C water into a 300-ml beaker, stirring the resulting aqueous slurry for 30 minutes, and then measuring the pH.

The referenced n value, the number average degree of condensation, may be determined by means of ^{31}P solid NMR. From J.R. van Wazer, C.F. Callis, J. Shoolery and R. Jones, J. Am. Chem. Soc., 78, 5715, 1956, the number of neighboring phosphate groups is known to give a unique 'chemical shift', which makes it possible to clearly distinguish between orthophosphates, pyrophosphates and polyphosphates.

Further, a process has been found for the preparation of the desired polyphosphate salt of a 1,3,5-triazine compound having an n value of at least 20, and preferably at least 40, and a M/P ratio of at least 1.1. This process involves the conversion of a 1,3,5-triazine compound with orthophosphoric acid into its orthophosphate salt, followed by dehydration and thermal treatment to convert the orthophosphate salt into a polyphosphate of the 1,3,5-triazine compound.

This thermal treatment is preferably performed at a temperature of at least 300°C, and preferably at least 310°C. In addition to orthophosphates of 1,3,5-triazine compounds, other 1,3,5-triazine phosphates may also be used, including, for example, a mixture of orthophosphates and pyrophosphates.

The orthophosphate of the 1,3,5-triazine compound may be prepared in a variety of processes. The preferred process involves adding the 1,3,5-triazine compound to an aqueous solution of orthophosphoric acid. An alternative process involves adding orthophosphoric acid to an aqueous slurry of the 1,3,5-triazine compound.

The process according to the present invention can also be carried out in the presence of a catalyst. As a result, the end product has better electrical properties as indicated by the Comparative Tracking Index (CTI) known from literature, measured according to the IEC 695-2-1 standard. Although any hydroxide may be utilized as a catalyst, alkali metal hydroxides and alkaline earth metal hydroxides are preferred. Salts of boric acid, for example zinc borate, may also be utilized as a catalyst. If used, the amount of catalyst used will generally be between 0.1 wt% and 10 wt%.

The reaction time required for satisfactory production of the desired polyphosphate of the 1,3,5-triazine compound is generally at least two minutes, and more generally at least five minutes, and generally less than 24 hours.

A polyphosphate of the 1,3,5-triazine derivative according to the present invention should contain less than 1 wt% of water-soluble material, and preferably less than 0.1 wt%. This low water-soluble

material content indicates that the product consists primarily of the desired polyphosphate.

It has also been found that polyphosphate salts of 1,3,5-triazine compounds according to the present invention are particularly suitable as flame retardants in polymer compositions. When used in this manner, the amount of flame retardant used in a polymer composition generally ranges from 15 to 45 wt%, and more generally from 20 to 40 wt%. It is believed that the suitability of these particular 1,3,5-triazine polyphosphate salts of compounds results from the increased thermal stability and increased pH achieved by compounds according to the present invention when compared with other flame retardants, such as halogen compounds, melamine, etc.

The flame retardant polymer compositions according to the present invention preferably comprise the following components:

- 35 - 55 wt% of polymer
- 20 15 - 45 wt% of polyphosphate salt of a 1,3,5-triazine compound with a number average degree of condensation n higher than 20
- 0 - 50 wt% of reinforcing fiber
- 0 - 20 wt% of carbon-forming compound
- 25 0 - 10 wt% of a catalyst promoting carbon formation

Suitable 1,3,5-triazine compounds include 2,4,6-triamine-1,3,5-triazine (melamine), melam, melem, melon, ammeline, ammelide, 2-ureidomelamine, acetoguanamine, benzoguanamine, diamine phenyltriazine or mixtures hereof. Melamine, melam, melem, melon or mixtures thereof, are preferred, and melamine in particular is preferred.

Polymers and polymer compositions to which polyphosphate salts of 1,3,5-triazine compounds

prepared according to the present invention may be added to improve flame retardant properties include the following:

- 5 1. Polymers of mono- and diolefins, for example polypropylene (PP), polyisobutylene, polybutylene-1, polymethylpentene-1, polyisoprene or polybutadiene; polyethylenes (optionally crosslinked) including, for example, high-density
10 polyethylene (HDPE), low-density polyethylene (LDPE), linear low-density polyethylene (LLDPE), or mixtures of these polymers.
2. Copolymers of mono- and diolefins, optionally including other vinyl monomers such as, for
15 example, ethylene-propylene copolymers, linear low-density polyethylene, and mixtures thereof with low-density polyethylene, as well as terpolymers of ethylene with propylene and a diene such as hexadiene, dicyclopentadiene or
20 ethylidene norbornene; furthermore, mixtures of such copolymers with the polymers listed under 1 such as, for example, polypropylene/ethylene-propylene copolymers.
3. Polystyrene, poly-(p-methyl-styrene), poly-(α -
25 methylstyrene) and copolymers of styrene or α -methylstyrene with dienes or acryl derivatives, such as, for example, styrene-butadiene, styrene-acrylonitrile, styrene-alkylmethacrylate, styrene-butadiene-alkylacrylate, styrene-maleic
30 anhydride and styrene-acrylonitrile-methylacrylate.
4. Polyphenylene oxide and polyphenylene sulphide and their mixtures with styrene polymers or with polyamides.

5. Polyurethanes derived from polyethers, polyesters and polybutadiene with terminal hydroxy groups on the one hand and aliphatic or aromatic polyisocyanates on the other, as well as their precursors.
6. Polyamides and copolyamides derived from diamines and dicarboxylic acids and/or from aminocarboxylic acids or the corresponding lactams, such as polyamide 4, polyamide 6, polyamide 6/6, 6/10, 6/9, 6/12, 4/6, 66/6, 6/66, polyamide 11, polyamide 12, aromatic polyamides based on an aromatic diamine and adipic acid; polyamides prepared from hexamethylene diamine and iso- and/or terephthalic acid and optionally an elastomer as modifier, for example poly-2,4,4-trimethyl hexamethylene terephthalamide, poly-m-phenylene-isophthalamide.
7. Polyesters derived from dicarboxylic acids and dialcohols and/or from hydroxycarboxylic acids or the corresponding lactones such as polyethylene terephthalate, polybutylene terephthalate, poly-1,4-dimethylol cyclohexane terephthalate and polyhydroxybenzoates.
8. Thermosetting resins including, for example, unsaturated polyesters, saturated polyesters, alkyd resins, polyacrylate or polyether or compositions containing one or more of these polymers and a crosslinking agent.

If reinforcing materials are used in a polymer composition according to the invention, their content may vary within wide limits depending in large part on the the desired or necessary level of one or more mechanical properties, as well as aesthetic, manufacturing, or economic considerations. In general, however, the amount of reinforcing materials will be

between 5 and 50 wt%, and more preferably, between 15 and 35 wt%. The reinforcing material can be chosen from the group of inorganic reinforcing materials such as, for example, mica, clay or glass fibers; or aramide
5 fibers and/or carbon fibers, or combinations thereof. In general, however, glass fibers are preferred.

The flame retardant action of the polyphosphate salt of a 1,3,5-triazine compound can be enhanced by the presence of a compound with a
10 synergistic effect for the flame retardant, particularly a so-called carbon-forming compound, optionally in combination with a catalyst promoting carbon formation. In general, the presence of a carbon-forming compound, with or without a catalyst, makes it
15 possible to reduce the triazine derivative polyphosphate content without reducing the flame retardant properties of the resulting polymer composition.

A number of substances are known to
20 reinforce the flame retardant action of the triazine derivative polyphosphate and may be included in the polymer composition as a carbon-forming compound. These substances include, for example, phenol resins, epoxy resins, melamine resins, alkyd resins, allyl resins,
25 unsaturated polyester resins, silicon resins, urethane resins, acrylate resins, starch, glucose, and compounds with at least two hydroxy groups. Examples of compounds with at least two hydroxy groups include various alcohols such as pentaerythritol, dipentaerythritol,
30 tripentaerythritol, and mixtures thereof. The concentration of such carbon-forming compounds in the polymer composition is typically less than 20 wt%, and preferably between 5 and 15 wt%.

A variety of catalysts may also be
35 incorporated to promote carbon formation. These

catalysts include, inter alia, metal salts of tungstic acid, complex acid oxides of tungsten with a metalloid, salts of tin oxide, ammonium sulphamate and/or its dimer. Metal salts of tungstic acid are preferably
5 alkali metal salts, and in particular sodium tungstate. A complex acid oxides of tungsten with a metalloid are understood to be complex acid oxides formed from a metalloid such as silicon or phosphorus and tungsten. The amount of catalyst used in the polymer composition
10 is generally 0.1 - 5 wt%, and preferably 0.1 - 2.5 wt%.

If polyolefins such as polyethylene, polypropylene, or mixtures thereof are used in the flame retardant polymer composition, it is preferred to also include a carbon-forming compound and/or a
15 catalyst for promoting carbon formation.

The flame retardant action of the polyphosphate salt of the 1,3,5-triazine compound can be further enhanced through the addition of a second flame retardant component. In principle any other known
20 flame retardant may be used as the second flame retardant component. Examples include antimony oxides, for example antimony trioxide; alkali earth metal oxides, for example magnesium oxide; other metal oxides, for example alumina, silica, zinc oxide, iron
25 oxide and manganese oxide; metal hydroxides, for example magnesium hydroxide and aluminium hydroxide; metal borates, for example hydrated or non-hydrated zinc borate; and phosphorus containing compounds. Examples of phosphorus containing compounds are zinc
30 phosphate, ammonium phosphate, ammonium pyrophosphate, ammonium polyphosphate, ethylene-diamine phosphate, piperazine phosphate, piperazine-pyrophosphate, melamine phosphate, dimelamine phosphate, melamine pyrophosphate, guanidine phosphate, dicyanodiamide
35 phosphate and/or urea phosphate. Phosphonates and

phosphate esters can also be used. Their content may vary within wide limits but generally does not exceed the content of the triazine derivative polyphosphate.

The polymer composition may further contain
5 the other customary additives, for example stabilizers, release agents, flow agents, dispersants, colorants and/or pigments, in amounts that are generally applicable. The additive content of the polymer compositions is generally selected to ensure that the
10 desired properties remain within acceptable limits, limits that will, of course, vary with the polymer composition and the intended application(s).

Polymer compositions according to the present invention can be prepared using most
15 conventional techniques, including the dry mixing of all or a number of components in a tumble mixer, followed by melting in a melt mixer, for example a Brabender mixer, a single-screw extruder, or, preferably, a twin-screw extruder.

20 The various components of the polymer composition of the invention can be fed together into the throat of the extruder or can be fed into the extruder singly or in sub-combinations through a plurality of inlets. If glass fiber reinforcement is to
25 be included in the composition, adding the glass fibers to the composition at the throat of the extruder is preferably avoided to minimize glass fiber breakage. A number of the components, for example colorants, stabilizers, and other additives, can be added to the
30 polymer as a masterbatch. The resulting polymer composition can then be processed into a variety of semi-manufactures and end products using a variety of techniques known to one skilled in the art, for example injection molding.

35 The invention will be elucidated with

reference to the following examples:

Comparative example:

A 50-liter reactor equipped with a stirrer
5 was charged with 29.25 l of pure water. While stirring,
8.619 kg of room temperature orthophosphoric acid (85
wt% H_3PO_4). Due to the exothermic reaction the
temperature of the diluted phosphoric acid solution
rose, and was kept at 50°C for 10 minutes. While still
10 stirring, 9.419 kg of melamine was then slowly added
(to prevent lump formation) to the solution. After the
melamine had been added, the reactor pressure was
reduced and the temperature increased to evaporate the
water and obtain a product with a moisture content of
15 less than 0.1 wt%. The resulting melamine phosphate,
with an M/P ratio of 1.0, was then heated to a
temperature of 310°C and converted into melamine
polyphosphate having an M/P ratio of 0.94. A 10 wt%
aqueous slurry of the resulting melamine polyphosphate
20 at 25°C had a pH less than 5.

A mixture consisting of 25 wt% of the
resulting melamine polyphosphate, 20 wt% of glass fiber
(PPG 3545 from PPG Industries) and 55 wt% of polyamide
6.6 (Durethan A31 from Bayer) prepared and extruded as
25 a granulate. Test bars were then prepared from the
resulting granulate and the following properties
determined:

Flame retardancy: V-1 according to UL-94 VB 1.6 mm.
Tensile strength: 140 MPa according to ISO 527
30 Elongation at break: 1.5% according to ISO 527
Charpy notch impact value: 37 KJ/m² according to ISO
179-1E-A/U
Modulus of elasticity: 11 GPa according to ISO 527.

Example:

A 50-liter reactor equipped with a stirrer was charged with 29.25 l of pure water. While stirring, 8.619 kg of room temperature orthophosphoric acid (85 wt% H_3PO_4) was added to the water. Due to the exothermic reaction the temperature of the diluted phosphoric acid solution rose and it was kept at 50°C for 10 minutes. While still stirring, 12.245 kg of melamine was then slowly added (to prevent lump formation). After the melamine had been added, the reactor pressure was reduced lowered and the temperature increased to evaporate the water and obtain a product with a moisture content of less than 0.1 wt%. The resulting melamine phosphate, having an M/P ratio of 1.3, was then heated to a temperature of 310°C and converted into melamine polyphosphate having an M/P ratio of 1.26. A 10 wt% aqueous slurry of the polyphosphate salt had a pH greater than 5.

A mixture consisting of 25 wt% of the resulting melamine polyphosphate according to the present invention, 20 wt% of glass fiber (PPG 3545 from PPG Industries) and 55 wt% of polyamide 6.6 (Durethan A31 from Bayer) prepared and extruded as a granulate. As with the polymer composition prepared in the comparative example, test bars were then made from the granulate and the following properties were determined:

Flame retardancy: V-0 according to UL-94 VB 1.6 mm.
tensile strength: 153 MPa according to ISO 527
elongation at break: 2.1% according to ISO 527
Charpy notch impact value: 48 KJ/m² according to ISO 179-1E-A/U
modulus of elasticity: 12 GPa according to ISO 527

CLAIMS

1. Polyphosphate salt of a 1,3,5-triazine compound,
characterized in that the number average degree
5 of condensation n is higher than 20 and with the
melamine content amounting to more than 1.1 mole
of melamine per mole of phosphorus atom.
2. Polyphosphate salt according to claim 1,
characterized in that the number average degree
10 of condensation n is higher than 40.
3. Polyphosphate salt of a 1,3,5-triazine compound,
characterized in that the melamine content is
higher than 1.2 moles of melamine per mole of
phosphorus atom.
- 15 4. Polyphosphate salt of a 1,3,5-triazine compound
according to any one of claims 1-3, characterized
in that the pH of a 10% slurry of the salt in
water is higher than or equal to 5.
5. Process for the preparation of the polyphosphate
20 salt of a 1,3,5-triazine compound with a number
average degree of condensation n that is higher
than 20 by converting a 1,3,5-triazine compound
with orthophosphoric acid at room temperature
into the phosphate of the 1,3,5-triazine
25 compound, after which this salt is converted into
the polyphosphate of the 1,3,5-triazine compound
via a thermal treatment.
6. Process according to claim 5, characterized in
that the phosphate of the 1,3,5-triazine compound
30 is converted into a polyphosphate of the 1,3,5-
triazine compound at a temperature higher than or
equal to 300°C.
7. Process according to claim 6, characterized in
that a number average degree of condensation

higher than 40 is obtained.

8. Process according to claims 5-7, characterized in that the phosphate of the 1,3,5-triazine compound is converted into the polyphosphate of the 1,3,5-triazine compound at a temperature higher than or equal to 310 °C.
9. Flame retardant polymer composition, characterized in that the polyphosphate of a 1,3,5-triazine compound with a number average degree of condensation of higher than 20 is used as flame retardant component.
10. Flame retardant polymer compositions comprising the following components:
 - 35 - 55 wt% of polymer
 - 15 - 45 wt% of polyphosphate salt of a 1,3,5-triazine compound with a number average degree of condensation n higher than 20
 - 0 - 50 wt% of reinforcing fiber
 - 0 - 20 wt% of carbon-forming compound
 - 0 - 10 wt% of catalyst promoting carbon formation
11. Flame retardant polymer composition according to claim 10, characterized in that melamine, melam, melem, melon or mixtures of these are used as 1,3,5-triazine compounds.
12. Flame retardant polymer composition according to claim 11, characterized in that melamine is used as 1,3,5-triazine compound.
13. Flame retardant polymer composition according to claims 9-12, characterized in that a flame retardant composition comprising the polyphosphate salt of a 1,3,5-triazine compound and a second flame retardant is used.

INTERNATIONAL SEARCH REPORT

International Application No
PC1/NL 99/00426

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07D251/54 C09K21/12 C08K5/3492 C08G79/04

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C07D C09K C08K C08G

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 97 44377 A (NISSAN CHEMICAL IND LTD ;SHINDO MASUO (JP); SHISHIDO KOUJI (JP); S) 27 November 1997 (1997-11-27) cited in the application see comparative example 1 claim 1	1,3,9
X	VOLFKOVIC ET AL.: "Condensed phosphates of melamine" Z. ANORG. ALLG. CHEM., no. 457, 1979, pages 20-30, XP002073286 see page 25/26 and 28 -/--	1,3,5

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

6 October 1999

Date of mailing of the international search report

13/10/1999

Name and mailing address of the ISA
European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

De Jong, B

INTERNATIONAL SEARCH REPORT

International Application No
PCT/NL 99/00426

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	CHEMICAL ABSTRACTS, vol. 112, no. 6, 5 February 1990 (1990-02-05) Columbus, Ohio, US; abstract no. 37343w, MUSZKO ET AL.: "Preparation of melamine polyphosphate" page 64; XP002073287 abstract & PL 143 704 A ----	1,3,9
X	WO 98 08898 A (DU PONT) 5 March 1998 (1998-03-05) page 7, line 22 - line 28; claim 1 ----	9
A	US 4 272 414 A (VANDERSALL H LAWRENCE) 9 June 1981 (1981-06-09) page 8, line 30; claim 1 ----	1,3,9
A	EP 0 015 006 A (BENCKISER KNAPSACK GMBH) 3 September 1980 (1980-09-03) example 8 ----	1,3
A	CA 1 032 941 A (ENCOAT CHEMICALS LIMITED) 13 June 1978 (1978-06-13) page 5 -page 6; example 1 ----	1,3,9
A	US 4 043 987 A (JOLICOEUR CARMEL REJEAN ET AL) 23 August 1977 (1977-08-23) example 1 -----	1,3,9

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/NL 99/00426

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9744377 A	27-11-1997	AU 2791397 A CA 2238474 A CN 1209824 A EP 0861283 A JP 10081691 A NO 983320 A	09-12-1997 27-11-1997 03-03-1999 02-09-1998 31-03-1998 17-07-1998
PL 143704 A		NONE	
WO 9808898 A	05-03-1998	AU 4088797 A EP 0922072 A	19-03-1998 16-06-1999
US 4272414 A	09-06-1981	AU 520286 B AU 4744379 A BR 7903262 A CA 1107047 A CH 641365 A DE 2921306 A FR 2426477 A GB 2021407 A,B GR 68366 A IT 1121505 B JP 1469645 C JP 54156397 A JP 63017114 B NZ 190550 A	21-01-1982 29-11-1979 11-12-1979 18-08-1981 29-02-1984 29-11-1979 21-12-1979 05-12-1979 23-12-1981 02-04-1986 14-12-1988 10-12-1979 12-04-1988 19-12-1980
EP 0015006 A	03-09-1980	DE 2907453 A AT 7684 T CA 1141143 A US 4293526 A US 4343779 A	04-09-1980 15-06-1984 15-02-1983 06-10-1981 10-08-1982
CA 1032941 A	13-06-1978	US 4043987 A	23-08-1977
US 4043987 A	23-08-1977	CA 1032941 A	13-06-1978